



NEWS RELEASE

PerkinElmer Agrees to Acquire Viral Vector Gene Delivery Leader SIRION Biotech

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Solutions to complement Company's growing position in cell and gene therapy research and development

WALTHAM, Mass.--(BUSINESS WIRE)--Jun. 22, 2021-- **PerkinElmer, Inc.** (NYSE: PKI) ("PerkinElmer") is pleased to announce it has entered into an agreement to acquire **SIRION Biotech GmbH**, a leading, global provider of viral vector-based technologies that drive improved delivery performance for cell and gene therapies. The acquisition is expected to close during the third quarter of 2021.

Headquartered in Munich, Germany, privately held SIRION has approximately 50 employees based in Germany, the U.S. and France. The company has established a strong licensing portfolio leveraged by over a dozen major pharmaceutical and biotech players researching more than twenty five diseases and conditions.

The addition of SIRION's offerings will complement **PerkinElmer's Horizon Discovery portfolio** which includes gene editing and modulation tools for CRISPR, CRISPRi and RNAi, custom cell lines for bio production and base editing technologies. Moreover, the acquisition will further broaden **PerkinElmer's existing cell and gene research solutions** featuring industry-leading high content, in vivo, and cell painting screening technologies; innovative immunoassays; cell plate readers; and advanced automation, microfluidics and informatics and analytical platforms.

Commenting on the agreement to acquire the company, Prahlad Singh, president and chief executive officer of

PerkinElmer, said, “Seventy percent of gene therapy trials today leverage viral vector approaches and we expect this to remain strong going forward given the demand for targeted, high-payload delivery in treating diseases like cancer. By combining SIRION’s innovative “payload” transport technology with our existing Horizon genetic material editing tools and phenotypic research solutions, we will be able to support organizations’ cell and gene therapy workflows as they look to streamline and accelerate their efforts.”

Dr. Christian Thirion, CEO of SIRION, added, “Our team is excited to work with PerkinElmer to continue to grow our leading position in viral vector technologies for cell and gene therapy. We will benefit greatly from becoming part of the PerkinElmer portfolio, with increased access to genomics analysis, gene editing and base editing technologies, as well as a strong global infrastructure and reach.”

To learn more about PerkinElmer’s full range of life sciences solutions, Informatics and OneSource services please visit: <https://www.perkinelmer.com/corporate/what-we-do/markets/life-sciences.html>.

About PerkinElmer

PerkinElmer enables scientists, researchers, and clinicians to address their most critical challenges across science and healthcare. With a mission focused on innovating for a healthier world, we deliver unique solutions to serve the diagnostics, life sciences, food, and applied markets. We strategically partner with customers to enable earlier and more accurate insights supported by deep market knowledge and technical expertise. Our dedicated team of about 14,000 employees worldwide is passionate about helping customers work to create healthier families, improve the quality of life, and sustain the well-being and longevity of people globally. The Company reported revenue of approximately \$3.8 billion in 2020, serves customers in 190 countries, and is a component of the S&P 500 index. Additional information is available through 1-877-PKI-NYSE, or at www.perkinelmer.com.

About SIRION Biotech GmbH

SIRION Biotech was founded in 2005 with the goal to spark a new generation of viral vector technologies for gene and cell therapy as well as vaccine development. SIRION evolves novel therapeutic viral vectors and uses proprietary technology platforms based on lenti-, adeno-, and adeno-associated viruses, to expedite its partners’ advances in drug development. For additional information, please visit www.sirion-biotech.com.

Factors Affecting Future Performance

This press release contains "forward-looking" statements within the meaning of the Private Securities Litigation Reform Act of 1995, including, but not limited to, statements relating to estimates and projections of future earnings per share, cash flow and revenue growth and other financial results, developments relating to our

customers and end-markets, and plans concerning business development opportunities, acquisitions and divestitures. Words such as "believes," "intends," "anticipates," "plans," "expects," "projects," "forecasts," "will" and similar expressions, and references to guidance, are intended to identify forward-looking statements. Such statements are based on management's current assumptions and expectations and no assurances can be given that our assumptions or expectations will prove to be correct. A number of important risk factors could cause actual results to differ materially from the results described, implied or projected in any forward-looking statements. These factors include, without limitation: (1) markets into which we sell our products declining or not growing as anticipated; (2) effect of the COVID-19 pandemic on our sales and operations; (3) fluctuations in the global economic and political environments; (4) our failure to introduce new products in a timely manner; (5) our ability to execute acquisitions and license technologies, or to successfully integrate acquired businesses and licensed technologies into our existing business or to make them profitable, or successfully divest businesses; (6) our failure to adequately protect our intellectual property; (7) the loss of any of our licenses or licensed rights; (8) our ability to compete effectively; (9) fluctuation in our quarterly operating results and our ability to adjust our operations to address unexpected changes; (10) significant disruption in third-party package delivery and import/export services or significant increases in prices for those services; (11) disruptions in the supply of raw materials and supplies; (12) the manufacture and sale of products exposing us to product liability claims; (13) our failure to maintain compliance with applicable government regulations; (14) regulatory changes; (15) our failure to comply with healthcare industry regulations; (16) economic, political and other risks associated with foreign operations; (17) our ability to retain key personnel; (18) significant disruption in our information technology systems, or cybercrime; (19) our ability to obtain future financing; (20) restrictions in our credit agreements; (21) discontinuation or replacement of LIBOR; (22) the United Kingdom's withdrawal from the European Union; (23) our ability to realize the full value of our intangible assets; (24) significant fluctuations in our stock price; (25) reduction or elimination of dividends on our common stock; and (26) other factors which we describe under the caption "Risk Factors" in our most recent quarterly report on Form 10-Q and in our other filings with the Securities and Exchange Commission. We disclaim any intention or obligation to update any forward-looking statements as a result of developments occurring after the date of this press release.

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